(FILE 'HOME' ENTERED AT 07:59:05 ON 31 JUL 2006)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 07:59:21 ON 31 JUL 2006

- L1 6 S (48 DIFFERENT ANTIBOD?)
- L2 2 S L1 AND ARRAY?
- L3 0 S L1 AND PD<2000
- L4 429 S (ANTIBOD? MICROARRAY)
- L5 3 S L4 AND PD<2000
- L6 214 DUPLICATE REMOVE L4 (215 DUPLICATES REMOVED)
- L7 35 S L6 AND REVIEW?

=>

(FILE 'HOME' ENTERED AT 07:59:05 ON 31 JUL 2006)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 07:59:21 ON 31 JUL 2006

L1	6	S	(48	DIFFERENT	ANTIBOD?)
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L2 2 S L1 AND ARRAY?

0 S L1 AND PD<2000

L4 429 S (ANTIBOD? MICROARRAY)

3 S L4 AND PD<2000

214 DUPLICATE REMOVE L4 (215 DUPLICATES REMOVED)

L7 35 S L6 AND REVIEW?

=>

L3

L5

L6

(FILE 'HOME' ENTERED AT 07:59:05 ON 31 JUL 2006)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 07:59:21 ON 31 JUL 2006

L1 6 S (	48	DIFFERENT	ANTIBOD?)
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2 S L1 AND ARRAY? L2L3

0 S L1 AND PD<2000

429 S (ANTIBOD? MICROARRAY) L4

3 S L4 AND PD<2000 L5

=>

(FILE 'HOME' ENTERED AT 07:59:05 ON 31 JUL 2006)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 07:59:21 ON 31 JUL 2006

L1 6 S (48 DIFFERENT ANTIBOD?)

L2 2 S L1 AND ARRAY?

0 S L1 AND PD<2000

L4 429 S (ANTIBOD? MICROARRAY)

L5 3 S L4 AND PD<2000

=>

L3

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ANSWER 18 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
     2004:951644 CAPLUS
DN
     142:293875
ED
     Entered STN: 10 Nov 2004
     High-throughput proteomics using antibody microarrays
ΤI
UA
     Wingren, Christer; Borrebaeck, Carl A. K.
     Department of Immunotechnology, Lund University, Lund, Swed.
CS
SO
     Expert Review of Proteomics (2004), 1(3), 355-364
     CODEN: ERPXA3; ISSN: 1478-9450
PB
     Future Drugs Ltd.
DT
     Journal; General Review
LA
     English
     9-0 (Biochemical Methods)
CC
     Section cross-reference(s): 1, 14
AB
     A review. Antibody-based microarrays are a novel technol. that
     hold great promise in proteomics. Microarrays can be printed with
     thousands of recombinant antibodies carrying the desired specificities,
     the biol. sample (e.g., an entire proteome) and any specifically bound
     analytes detected. The microarray patterns that are generated can then be
     converted into proteomic maps, or mol. fingerprints, revealing the composition
     of the proteome. Using this tool, global proteome anal. and protein
     expression profiling will thus provide new opportunities for biomarker
     discovery, drug target identification and disease diagnostics, as well as
     providing insights into disease biol. Intense work is currently underway
     to develop this novel technol. platform into the high-throughput proteomic
     tool required by the research community.
ST
     review high throughput proteomic antibody
     microarray
IT
     Protein expression profiles
     Protein microarray technology
     Proteomics
        (high-throughput proteomics using antibody
        microarrays)
IT
     Proteome
     RL: ANT (Analyte); ANST (Analytical study)
        (high-throughput proteomics using antibody
        microarrays)
IT
     Antibodies and Immunoglobulins
     RL: ARG (Analytical reagent use); DEV (Device component use); ANST
     (Analytical study); USES (Uses)
        (high-throughput proteomics using antibody
        microarrays)
RE.CNT
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ANSWER 2 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN 2005:219591 BIOSIS AN PREV200510003108 DN Progress in protein and antibody microarray ТT technology. Angenendt, Philipp [Reprint Author] AU German Canc Res Ctr, Funct Genome Anal, Neuenheimer Feld 580, D-69120 CS Heidelberg, Germany p.angenendt@dkfz-heidelberg.de Drug Discovery Today, (APR 1 2005) Vol. 10, No. 7, pp. 503-511. SO ISSN: 1359-6446. DTArticle General Review; (Literature Review) LA English Entered STN: 10 Jun 2005 ED Last Updated on STN: 10 Jun 2005 AB The success of genome sequencing projects has led to a shift from the description of single molecules to the characterisation of complex samples. At the same time, there is growing interest not only in studying organisms at the genomic level, but in the characterization of their proteome. Such a task would not be possible without the availability of appropriate technologies. Protein and antibody microarray technologies are, in addition to two-dimensional gel electrophoresis followed by mass spectrometry, two of the most propitious technologies for the screening of complex protein samples. Nevertheless, to succeed, protein and antibody microarrays have to overcome their current limitations. This review aims to introduce these new technologies and highlights their current prospects and limitations. CC Genetics - General 03502 Genetics - Population genetics 03509 Major Concepts IT Methods and Techniques; Population Genetics (Population Studies);

Molecular Genetics (Biochemistry and Molecular Biophysics)

IT Methods & Equipment

> mass spectrometry: laboratory techniques, spectrum analysis techniques; two-dimensional gel electrophoresis: electrophoretic techniques, laboratory techniques; genome sequencing: laboratory techniques, genetic techniques; antibody microarray: laboratory techniques; protein microarray: laboratory techniques

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ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
     1997:101881 CAPLUS
DN
     126:209077
ED
     Entered STN: 13 Feb 1997
     Microarray-based immunoassays
ΤI
     Chu, F. W.; Edwards, P. R.; Ekins, R. P.; Berger, H.; Finckh, P.; Krause,
AU
     Div. Mol. Endocrinology, Univ. College London Medical School, London, W1N
CS
     ACS Symposium Series (1997), 657 (Immunochemical Technology for
SO
     Environmental Applications), 170-184
     CODEN: ACSMC8; ISSN: 0097-6156
     American Chemical Society
PΒ
DT
     Journal; General Review
LA
     English
     9-0 (Biochemical Methods)
CC
     Section cross-reference(s): 3, 4
AB
     A review with 19 refs. about the general principles underlying the
     emerging technol. of microarray-based immunoassays. Recent worldwide
     interest in the development of miniaturized, array-based, multianalyte
     binding assay methods suggests that the ligand assay field is on the brink
     of a technol. revolution. Our own collaborative studies in this area have
     centered largely (but not exclusively) on antibody spot "immunoarrays"
     localized on "microchips" which are potentially capable of determining the
amts.
     of hundreds of different analytes in a small sample (such as a single drop
     of blood). Analogous technol. for genetic testing using oligonucleotide
     arrays is under active development both in the US and Europe. Array-based
     immunoassay methods are clearly likely to prove of particular importance
     in areas such as environmental monitoring where the concns. of many
     different analytes in test samples are required to be simultaneously determined
ST
     review microarray based immunoassay antibody
ΙT
     Immunoassay
        (apparatus; microarray-based immunoassays)
IT
     Blood analysis
     Immunoassay
        (microarray-based immunoassays)
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RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)

IT

Antibodies

(microarray-based immunoassays)

ANSWER 1 OF 3 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN 1999:301008 BIOSIS DNPREV199900301008 ΤI Immunoassay and other ligand assays: From isotopes to luminescence. Ekins, Roger [Reprint author] ΑU Department of Molecular Endocrinology, University College London Medical CS School, Mortimer Street, London, W1N 8AA, UK Journal of Clinical Ligand Assay, (Spring, 1999) Vol. 22, No. 1, SO pp. 61-77. print. ISSN: 1081-1672. DT Article General Review; (Literature Review) LA English ED Entered STN: 12 Aug 1999 Last Updated on STN: 12 Aug 1999 This article reviews key developments in the ligand assay field. These AB methods are represented by three generations: 1) competitive assays which rely on radio-labeled analyte markers to reveal the products of the reaction between the analyte and a specific binding agent; 2) non-competitive ultra-sensitive assays in which the binding agent is labeled with a non-isotopic marker of much higher specific activity than radioisotopes; and 3) microarray technologies which permit simultaneous ultra-sensitive measurement of tens, hundreds, or thousands of analytes in a sample. The importance of liqund assays stems largely from their high sensitivity. However, developments in the field have been profoundly affected by misunderstanding the concept of sensitivity itself. obscured the significance of major innovations such as in vitro methods of (monoclonal) antibody production and the microspot assay techniques that underlie microarray-based methods. These may revolutionize in vitro diagnostics in the next decade. Biochemistry methods - General CC 10050 General biology - Information, documentation, retrieval and computer 00530 applications Biochemistry studies - General 10060 Immunology - General and methods 34502 ITMajor Concepts Computer Applications (Computational Biology); Immune System (Chemical Coordination and Homeostasis); Methods and Techniques IT Methods & Equipment antibody microarray: immunologic method; immunoassay automation: computer method, immunologic method; ligand assay: analytical method; microanalytical chips: computer system;

ultra-sensitive multianalyte assay: analytical method; DNA analysis:

analytical method

luminescence

Miscellaneous Descriptors

IT

ANSWER 1 OF 3 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN AN1999:301008 BIOSIS PREV199900301008 DN Immunoassay and other ligand assays: From isotopes to luminescence. ΤI Ekins, Roger [Reprint author] ΑŲ Department of Molecular Endocrinology, University College London Medical CS School, Mortimer Street, London, W1N 8AA, UK Journal of Clinical Ligand Assay, (Spring, 1999) Vol. 22, No. 1, SO pp. 61-77. print. ISSN: 1081-1672. DTArticle General Review; (Literature Review) LA Entered STN: 12 Aug 1999 Last Updated on STN: 12 Aug 1999 This article reviews key developments in the ligand assay field. These AB methods are represented by three generations: 1) competitive assays which rely on radio-labeled analyte markers to reveal the products of the reaction between the analyte and a specific binding agent; 2) non-competitive ultra-sensitive assays in which the binding agent is labeled with a non-isotopic marker of much higher specific activity than radioisotopes; and 3) microarray technologies which permit simultaneous ultra-sensitive measurement of tens, hundreds, or thousands of analytes in a sample. The importance of liqand assays stems largely from their high sensitivity. However, developments in the field have been profoundly affected by misunderstanding the concept of sensitivity itself. This has obscured the significance of major innovations such as in vitro methods of (monoclonal) antibody production and the microspot assay techniques that underlie microarray-based methods. These may revolutionize in vitro diagnostics in the next decade. CC Biochemistry methods - General 10050 General biology - Information, documentation, retrieval and computer 00530 applications Biochemistry studies - General 10060 Immunology - General and methods 34502 IT Major Concepts Computer Applications (Computational Biology); Immune System (Chemical

Coordination and Homeostasis); Methods and Techniques

ITMethods & Equipment

antibody microarray: immunologic method; immunoassay automation: computer method, immunologic method; ligand assay: analytical method; microanalytical chips: computer system; ultra-sensitive multianalyte assay: analytical method; DNA analysis: analytical method

ITMiscellaneous Descriptors luminescence

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ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
     1997:101881 CAPLUS
AN
     126:209077
DN
     Entered STN: 13 Feb 1997
ED
ΤI
    Microarray-based immunoassays
     Chu, F. W.; Edwards, P. R.; Ekins, R. P.; Berger, H.; Finckh, P.; Krause,
ΑU
    Div. Mol. Endocrinology, Univ. College London Medical School, London, W1N
CS
    ACS Symposium Series (1997), 657 (Immunochemical Technology for
SO
    Environmental Applications), 170-184
     CODEN: ACSMC8; ISSN: 0097-6156
PΒ
    American Chemical Society
     Journal; General Review
DT
     English
LA
     9-0 (Biochemical Methods)
CC
     Section cross-reference(s): 3, 4
    A review with 19 refs. about the general principles underlying the
AB
     emerging technol. of microarray-based immunoassays. Recent worldwide
     interest in the development of miniaturized, array-based, multianalyte
    binding assay methods suggests that the ligand assay field is on the brink
     of a technol. revolution. Our own collaborative studies in this area have
     centered largely (but not exclusively) on antibody spot "immunoarrays"
     localized on "microchips" which are potentially capable of determining the
amts.
     of hundreds of different analytes in a small sample (such as a single drop
     of blood). Analogous technol. for genetic testing using oligonucleotide
     arrays is under active development both in the US and Europe. Array-based
     immunoassay methods are clearly likely to prove of particular importance
     in areas such as environmental monitoring where the concns. of many
    different analytes in test samples are required to be simultaneously determined
     review microarray based immunoassay antibody
ST
IT
    Immunoassay
        (apparatus; microarray-based immunoassays)
IT
    Blood analysis
    Immunoassay
        (microarray-based immunoassays)
```

RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)

IT

Antibodies

(microarray-based immunoassays)

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ANSWER 31 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
    2001:182232 CAPLUS
DN
     135:32439
ED
     Entered STN: 16 Mar 2001
     Protein and antibody arrays and their medical applications
TΤ
    Cahill, D. J.
ΑU
    Max-Planck-Institute of Molecular Genetics, Berlin, D-14195, Germany
CS
     Journal of Immunological Methods (2001), 250(1-2), 81-91
SO
     CODEN: JIMMBG; ISSN: 0022-1759
    Elsevier Science B.V.
PB
     Journal; General Review
DT
LA
    English
CC
     15-0 (Immunochemistry)
     Section cross-reference(s): 9, 14
    A review with 30 refs. Many new gene products are being
AB
     discovered by large-scale genomics and proteomics strategies, the
     challenge is now to develop high throughput approaches to systematically
     analyze these proteins and to assign a biol. function to them. Having
     access to these gene products as recombinantly expressed proteins, would
     allow them to be robotically arrayed to generate protein chips. Other
     applications include using these proteins for the generation of specific
     antibodies, which can also be arrayed to produce antibody chips. The
     availability of such protein and antibody arrays would facilitate the
     simultaneous anal. of thousands of interactions within a single experiment
     This chapter will focus on current strategies used to generate protein and
     antibody arrays and their current applications in biol. research, medicine
     and diagnostics. The shortcomings of these approaches, the developments
     required, as well as the potential applications of protein and antibody
     arrays will be discussed.
     review protein antibody microarray diagnosis
ST
IT
    Antibodies
     RL: ANT (Analyte); ARG (Analytical reagent use); THU (Therapeutic use);
     ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (protein and antibody arrays and their medical applications)
     Proteins, general, biological studies
TT
     RL: ANT (Analyte); THU (Therapeutic use); ANST (Analytical study); BIOL
     (Biological study); USES (Uses)
        (protein and antibody arrays and their medical applications)
IT
    Diagnosis
        (protein and antibody arrays in)
             THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
       30
RE
(1) Anderson, L; Electrophoresis 1997, V18, P533 CAPLUS
(2) Bussow, K; Nucleic Acids Res 1998, V26, P5007 CAPLUS
(3) Cahill, D; Proteomics From Protein Sequence To Function 2000, P1
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(8) Engelbert, C; Cancer Res 2000, V60, P1526
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ANSWER 31 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
     2001:182232 CAPLUS
DN
     135:32439
ED
     Entered STN: 16 Mar 2001
     Protein and antibody arrays and their medical applications
ΤI
ΑU
     Cahill, D. J.
     Max-Planck-Institute of Molecular Genetics, Berlin, D-14195, Germany
CS
SO
     Journal of Immunological Methods (2001), 250(1-2), 81-91
     CODEN: JIMMBG; ISSN: 0022-1759
     Elsevier Science B.V.
PB
     Journal; General Review
DT
LA
     English
     15-0 (Immunochemistry)
CC
     Section cross-reference(s): 9, 14
     A review with 30 refs. Many new gene products are being
     discovered by large-scale genomics and proteomics strategies, the
     challenge is now to develop high throughput approaches to systematically
     analyze these proteins and to assign a biol. function to them. Having
     access to these gene products as recombinantly expressed proteins, would
     allow them to be robotically arrayed to generate protein chips. Other
     applications include using these proteins for the generation of specific
     antibodies, which can also be arrayed to produce antibody chips. The
     availability of such protein and antibody arrays would facilitate the
     simultaneous anal. of thousands of interactions within a single experiment
     This chapter will focus on current strategies used to generate protein and
     antibody arrays and their current applications in biol. research, medicine
     and diagnostics. The shortcomings of these approaches, the developments
     required, as well as the potential applications of protein and antibody
     arrays will be discussed.
     review protein antibody microarray diagnosis
ST
IT
     Antibodies
     RL: ANT (Analyte); ARG (Analytical reagent use); THU (Therapeutic use);
     ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (protein and antibody arrays and their medical applications)
TT
     Proteins, general, biological studies
     RL: ANT (Analyte); THU (Therapeutic use); ANST (Analytical study); BIOL
     (Biological study); USES (Uses)
        (protein and antibody arrays and their medical applications)
IT
     Diagnosis
        (protein and antibody arrays in)
              THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
        30
RE
(1) Anderson, L; Electrophoresis 1997, V18, P533 CAPLUS
(2) Bussow, K; Nucleic Acids Res 1998, V26, P5007 CAPLUS
(3) Cahill, D; Proteomics From Protein Sequence To Function 2000, P1
(4) Cohen, C; Anal Biochem 1999, V273, P89 CAPLUS
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(6) Ekins, R; Clin Chem 1998, V44, P2015 CAPLUS
(7) Emili, A; Nat Biotechnol 2000, V18, P393 CAPLUS (8) Engelbert, C; Cancer Res 2000, V60, P1526 (9) Ge, H; Nucleic Acids Res 2000, V28, Pe3 CAPLUS
(10) Herwig, R; Genome Res 1999, V9, P1093 CAPLUS
(11) Holt, L; Nucleic Acids Res 2000, V28(1-5), Pe72
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     2002:787438 CAPLUS
     138:37532
     Entered STN: 16 Oct 2002
ED
     Antibody arrays: an embryonic but rapidly growing technology
TT
ΔIJ
     Lal, Sean P.; Christopherson, Richard I.; dos Remedios, Cristobal G.
     Institute for Biomedical Research, University of Sydney, Sydney, 2006,
CS
     Australia
SO
     Drug Discovery Today (2002), 7(18, Suppl.), S143-S149
     CODEN: DDTOFS; ISSN: 1359-6446
PΒ
     Elsevier Science Ltd.
DT
     Journal; General Review
     English
LA
CC
     15-0 (Immunochemistry)
     Section cross-reference(s): 9
     A review. Protein arrays are now an attractive proposition as
AB
     they can measure a diverse range of protein interactions not possible with
     traditional DNA arrays. Antibody arrays are a specific subset of this
     technol. Originally conceived as multi-analyte detectors, antibody arrays
     are now used in a wide variety of applications. For instance, the
     potential of this technol. to diagnose human diseases, such as leukemia,
     breast cancer and, potentially, heart failure, has stimulated much
     interest. Furthermore, identification of new protein targets in
     particular disease states will prove to be an invaluable tool in drug
     discovery and development. Patient prognosis and treatment are also
     potential applications of the technol. Antibody arrays have proved to be
     dynamic in response to these broad range of possibilities. This
     review examines variations in antibody array design and discusses
     current and potential applications of this novel and interesting technol.
ST
     review antibody microarray protein
ΙT
     Protein microarray technology
        (design and applications of antibody arrays)
IT
     Antibodies and Immunoglobulins
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (design and applications of antibody arrays)
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ANSWER 23 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
    2003:928473 CAPLUS
AN
DN
    140:406306
    Entered STN: 28 Nov 2003
ED
    Protein microarrays: A literature survey
ΤI
    Kricka, Larry J.; Joos, Thomas; Fortina, Paolo
ΑIJ
    Department of Pathology and Laboratory Medicine, 7.103 Founders Pavilion,
CS
    Medical Center, University of Pennsylvania, Philadelphia, PA, 19104, USA
    Clinical Chemistry (Washington, DC, United States) (2003), 49(12), 2109
    CODEN: CLCHAU; ISSN: 0009-9147
PΒ
    American Association for Clinical Chemistry
    Journal
\mathtt{DT}
LA
    20-5 (History, Education, and Documentation)
CC
    The Working Group has now completed a survey on the protein microarray
AB
     literature. The current survey covers the protein, peptide, and
    antibody, microarray literature up to the middle of
     2003. The literature survey has been divided into four sections: (1)
    General (books, reviews, editorials); (2) Fabrication (array
    construction and detection methodologies); (3) Applications (protein
     identification, and quantification, array-based proteomics, protein
    interactions); and (4) Patents (only US patents listed currently).
    protein microarray literature
ST
IT
    Databases
    Information systems
    Internet
    Literature
    Protein microarray technology
        (protein microarrays: a literature survey)
TТ
    Proteins
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (protein microarrays: a literature survey)
             THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
(1) Kricka, L; Clin Chem 2001, V47, P1479 CAPLUS
(2) Kricka, L; Clin Chem 2002, V48, P1620 CAPLUS
(3) Kricka, L; Clin Chem 2002, V48, P662 CAPLUS
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ANSWER 23 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
     2003:928473 CAPLUS
AN
DN
     140:406306
     Entered STN: 28 Nov 2003
ED
     Protein microarrays: A literature survey
ΤI
     Kricka, Larry J.; Joos, Thomas; Fortina, Paolo
AU
     Department of Pathology and Laboratory Medicine, 7.103 Founders Pavilion,
CS
     Medical Center, University of Pennsylvania, Philadelphia, PA, 19104, USA
     Clinical Chemistry (Washington, DC, United States) (2003), 49(12), 2109
SO
     CODEN: CLCHAU; ISSN: 0009-9147
PΒ
     American Association for Clinical Chemistry
     Journal
DT
LA
     English
     20-5 (History, Education, and Documentation)
CC
     The Working Group has now completed a survey on the protein microarray
AΒ
     literature. The current survey covers the protein, peptide, and
     antibody, microarray literature up to the middle of
           The literature survey has been divided into four sections: (1)
     General (books, reviews, editorials); (2) Fabrication (array
     construction and detection methodologies); (3) Applications (protein
     identification, and quantification, array-based proteomics, protein
     interactions); and (4) Patents (only US patents listed currently).
     protein microarray literature
ST
    Databases
TТ
     Information systems
     Internet
     Literature
     Protein microarray technology
        (protein microarrays: a literature survey)
IT
     Proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (protein microarrays: a literature survey)
              THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
(1) Kricka, L; Clin Chem 2001, V47, P1479 CAPLUS
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(3) Kricka, L; Clin Chem 2002, V48, P662 CAPLUS
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ANSWER 21 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
AN
     2004:375171 CAPLUS
DN
     141:309761
ED
     Entered STN: 10 May 2004
     Multiplexed protein analysis using spotted antibody
TI
     microarrays
ΑU
     Haab, Brian B.; Zhou, Heping
CS
     Van Andel Research Institute, Grand Rapids, MI, USA
SO
     Methods in Molecular Biology (Totowa, NJ, United States) (2004),
     264 (Protein Arrays), 33-45
     CODEN: MMBIED; ISSN: 1064-3745
PB
     Humana Press Inc.
     Journal; General Review
DT
LA
     English
CC
     9-0 (Biochemical Methods)
AB
     A review. This chapter describes methods for the production and use
     of antibody microarrays. The methods are divided into
     (a) antibody handling and microarray production, (b) sample preparation, and
(c)
     microarray use. Two types of detection methods are described: direct .
     labeling and a fluorescence-linked immunosorbent assay (FLISA). In the
     direct labeling method, all proteins in a complex mixture are labeled with
     either a fluorophore or a hapten that allows subsequent detection.
     FLISA detection, a capture antibody on the microarray captures the
     unlabeled protein target, which is detected by a detection antibody and a
     fluorophore-labeled secondary antibody. Each method has particular
     optimal uses, which are discussed in the text.
ST
     review multiplexed protein analysis spotted antibody
     microarrays fluorescence immunoassay
     Fluorescence immunoassay
TΤ
        (fluorescence-linked immunosorbent assay, FLISA; multiplexed protein
        anal. using spotted antibody microarrays)
TT
     Fluorescent indicators
     Sample preparation
        (multiplexed protein anal. using spotted antibody
        microarrays)
TT
     Proteins
     RL: ANT (Analyte); ANST (Analytical study)
        (multiplexed protein anal. using spotted antibody
        microarrays)
IT
     Antibodies and Immunoglobulins
     RL: ANT (Analyte); ARG (Analytical reagent use); BSU (Biological study,
     unclassified); ANST (Analytical study); BIOL (Biological study); USES
     (Uses)
        (multiplexed protein anal. using spotted antibody
        microarrays)
IT
     Blood analysis
        (serum profiling; multiplexed protein anal. using spotted
        antibody microarrays)
IT
     Protein microarray technology
        (spotted antibody; multiplexed protein anal. using spotted
        antibody microarrays)
              THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
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ANSWER 18 OF 35 CAPLUS COPYRIGHT 2006 ACS on STN
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DN
     142:293875
ED
     Entered STN: 10 Nov 2004
     High-throughput proteomics using antibody microarrays
TI
     Wingren, Christer; Borrebaeck, Carl A. K.
CS
     Department of Immunotechnology, Lund University, Lund, Swed.
     Expert Review of Proteomics (2004), 1(3), 355-364
SO
     CODEN: ERPXA3; ISSN: 1478-9450
     Future Drugs Ltd.
PB
     Journal; General Review
DT
     English
LA
     9-0 (Biochemical Methods)
CC
     Section cross-reference(s): 1, 14
     A review. Antibody-based microarrays are a novel technol. that
AB
     hold great promise in proteomics. Microarrays can be printed with
     thousands of recombinant antibodies carrying the desired specificities,
     the biol. sample (e.g., an entire proteome) and any specifically bound
     analytes detected. The microarray patterns that are generated can then be
     converted into proteomic maps, or mol. fingerprints, revealing the composition
     of the proteome. Using this tool, global proteome anal. and protein
     expression profiling will thus provide new opportunities for biomarker
     discovery, drug target identification and disease diagnostics, as well as
     providing insights into disease biol. Intense work is currently underway
     to develop this novel technol. platform into the high-throughput proteomic
     tool required by the research community.
ST
     review high throughput proteomic antibody
     microarray
IT
     Protein expression profiles
     Protein microarray technology
     Proteomics
        (high-throughput proteomics using antibody
        microarrays)
IT
     Proteome
     RL: ANT (Analyte); ANST (Analytical study)
        (high-throughput proteomics using antibody
        microarrays)
     Antibodies and Immunoglobulins
IT
     RL: ARG (Analytical reagent use); DEV (Device component use); ANST
     (Analytical study); USES (Uses)
        (high-throughput proteomics using antibody
        microarrays)
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ANSWER 2 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN AN 2005:219591 BIOSIS PREV200510003108 DN Progress in protein and antibody microarray тT technology. Angenendt, Philipp [Reprint Author] ΑU German Canc Res Ctr, Funct Genome Anal, Neuenheimer Feld 580, D-69120 CS Heidelberg, Germany p.angenendt@dkfz-heidelberg.de Drug Discovery Today, (APR 1 2005) Vol. 10, No. 7, pp. 503-511. SO ISSN: 1359-6446. DTArticle General Review; (Literature Review) LA English Entered STN: 10 Jun 2005 ED Last Updated on STN: 10 Jun 2005 The success of genome sequencing projects has led to a shift from the AB description of single molecules to the characterisation of complex samples. At the same time, there is growing interest not only in studying organisms at the genomic level, but in the characterization of their proteome. Such a task would not be possible without the availability of appropriate technologies. Protein and antibody microarray technologies are, in addition to two-dimensional gel electrophoresis followed by mass spectrometry, two of the most propitious technologies for the screening of complex protein samples. Nevertheless, to succeed, protein and antibody microarrays have to overcome their current limitations. This review aims to introduce these new technologies and highlights their current prospects and limitations. CC Genetics - General 03502 Genetics - Population genetics 03509 IT Major Concepts Methods and Techniques; Population Genetics (Population Studies); Molecular Genetics (Biochemistry and Molecular Biophysics)

TT Methods & Equipment

mass spectrometry: laboratory techniques, spectrum analysis techniques; two-dimensional gel electrophoresis: electrophoretic techniques, laboratory techniques; genome sequencing: laboratory techniques, genetic techniques; antibody microarray: laboratory techniques; protein microarray: laboratory techniques